

=> s Kontsekova E?/au
L1 48 KONTSEKOVA E?/AU

=> s l1 and tau
87233 TAU
168 TAUS
87285 TAU
(TAU OR TAUS)
L2 10 L1 AND TAU

=> s l2 and truncated
41439 TRUNCATED
L3 4 L2 AND TRUNCATED

=> d ibib abs 1-4

L3 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:607535 CAPLUS

DOCUMENT NUMBER: 145:121973

TITLE: Truncated tau from sporadic
Alzheimer's disease suffices to drive neurofibrillary
degeneration in vivo

AUTHOR(S): Zilka, Norbert; Filipcik, Peter; Koson, Peter;
Fialova, Lubica; Skrabana, Rostislav; Zilkova, Monika;
Rolkova, Gabriela; Kontsekova, Eva; Novak,
Michal

CORPORATE SOURCE: Axon Neuroscience GmbH, Vienna, 1030, Austria

SOURCE: FEBS Letters (2006), 580(15), 3582-3588

CODEN: FEBLAL; ISSN: 0014-5793

PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Truncated tau protein is the characteristic feature of
human sporadic Alzheimer's disease. We have identified truncated
tau proteins conformationally different from normal healthy
tau. Subpopulations of these structurally different tau
species promoted abnormal microtubule assembly in vitro suggesting toxic
gain of function. To validate pathol. activity in vivo we expressed
active form of human truncated tau protein as
transgene, in the rat brain. Its neuronal expression led to the
development of the neurofibrillary degeneration of Alzheimer's type.
Furthermore, biochem. anal. of neurofibrillary changes revealed that
massive sarcosyl insol. tau complexes consisted of human
Alzheimer's tau and endogenous rat tau in ratio 1:1
including characteristic Alzheimer's disease (AD)-specific proteins (A68).
This work represents first insight into the possible causative role of
truncated tau in AD neurofibrillary degeneration in
vivo.

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:60551 CAPLUS

DOCUMENT NUMBER: 140:124832

TITLE: Truncated tau proteins

INVENTOR(S): Kontsekova, Eva

PATENT ASSIGNEE(S): Axon Neuroscience Forschungs- und Entwicklungs GmbH,
Austria

SOURCE: PCT Int. Appl., 97 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004007547	A2	20040122	WO 2003-EP7389	20030709
WO 2004007547	A3	20040722		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2003253044	A1	20040202	AU 2003-253044	20030709
EP 1521774	A2	20050413	EP 2003-763763	20030709
EP 1521774	B1	20080827		
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
CN 1668641	A	20050914	CN 2003-816647	20030709
JP 2006515270	T	20060525	JP 2004-520541	20030709
AT 391781	T	20080415	AT 2003-763764	20030709
AT 406383	T	20080915	AT 2003-763763	20030709
ES 2304146	T3	20080916	ES 2003-763764	20030709
EP 1995255	A1	20081126	EP 2008-14706	20030709
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LI, LU, MC, NL, PT, RO, SE, SI, SK, TR, LT, LV			
ES 2311734	T3	20090216	ES 2003-763763	20030709
US 20060167227	A1	20060727	US 2005-521140	20051031
PRIORITY APPLN. INFO.:			AT 2002-1053	A 20020712
			EP 2003-763763	A3 20030709
			WO 2003-EP7389	W 20030709

AB Described are novel N- and C-terminally double truncated tau mols., (type IA, IB, IIA and IIB tau mols.) as well as methods for providing these mols., both from recombinant and biol. sources. Moreover, screening methods using these mols. in connection with Alzheimer's diagnosis and therapy are provided.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2003:64022 CAPLUS

DOCUMENT NUMBER: 138:367296

TITLE: DC11: a novel monoclonal antibody revealing Alzheimer's disease-specific tau epitope

AUTHOR(S): Vechterova, Lubica; Kontsekova, Eva; Zilka, Norbert; Ferencik, Miroslav; Ravid, Rivka; Novak, Michal

CORPORATE SOURCE: Axon Neuroscience, Vienna, A-1030, Austria

SOURCE: NeuroReport (2003), 14(1), 87-91

CODEN: NERPEZ; ISSN: 0959-4965

PUBLISHER: Lippincott Williams & Wilkins

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Using tau protein exts. from Alzheimer's disease (AD) brain tissue, we generated a monoclonal antibody (mAb DC11) which decorated

neurofibrillary pathol. in brain derived from AD patients on immunohistochem., and which lacked reactivity with healthy brain tissue. The same pattern of DC11 specificity was observed on Western blot. The main constituent of structures decorated by DC11 is microtubule-associated protein tau. In Western blot, DC11 recognized neither native healthy tau nor its full length recombinant counterpart. However, the mAb showed strong immunoreactivity with truncated tau (residues .tau.151-421), thus indicating the requirement for a conformational epitope. Importantly, the DC11 epitope was phosphorylation independent. The immunochem. parameters of mAb show that DC11 could represent a novel structural probe with the specificity for conformation of pathol. tau present in AD brains.

REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1995:814757 CAPLUS

DOCUMENT NUMBER: 123:250505

ORIGINAL REFERENCE NO.: 123:44639a,44642a

TITLE: Quick purification of recombinant human truncated tau proteins for immunoanalysis

AUTHOR(S): Kontsekkova, Eva; Cattaneo, Antonino; Novak, Michal

CORPORATE SOURCE: Institute of Virology, Slovak Academy of Sciences, 842 46, Bratislava, Czech.

SOURCE: Journal of Immunological Methods (1995), 185(2), 245-8
CODEN: JIMMBG; ISSN: 0022-1759

PUBLISHER: Elsevier

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A simple and rapid purification method is described which exploits the heat stability of human tau (.tau.) protein to prepare truncated forms of this protein derived from bacteria. Bacterial cells expressing .tau. fragments were pelleted, resuspended in phosphate buffered saline and boiled for 5 min. After centrifugation the supernatant containing thermostable .tau. was filtered (0.45 µm) and used for immunoanal. with monoclonal antibodies. The purified .tau. fragments exhibited identical antigenic properties as fragments isolated by a conventional procedure, based on ion exchange chromatog. on phosphocellulose. In contrast to the conventional approach, our method is less complicated, cheaper and significantly reduces the time required for isolation of the recombinant .tau. fragments.

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E1 THROUGH E27 ASSIGNED

=> d all 13 2

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

AN 2004:60551 CAPLUS

DN 140:124832

ED Entered STN: 26 Jan 2004

TI Truncated tau proteins

IN Kontsekkova, Eva

PA Axon Neuroscience Forschungs- und Entwicklungs GmbH, Austria

SO PCT Int. Appl., 97 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM C07K014-47
ICS C12N015-12; G01N033-53; A01K067-027; A61K039-00; C12P021-02;
C07K016-00

CC 9-2 (Biochemical Methods)
Section cross-reference(s): 3, 6, 13, 14, 15

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004007547	A2	20040122	WO 2003-EP7389	20030709
	WO 2004007547	A3	20040722		
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	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	AU 2003253044	A1	20040202	AU 2003-253044	20030709
	EP 1521774	A2	20050413	EP 2003-763763	20030709
	EP 1521774	B1	20080827		
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
	CN 1668641	A	20050914	CN 2003-816647	20030709
	JP 2006515270	T	20060525	JP 2004-520541	20030709
	AT 391781	T	20080415	AT 2003-763764	20030709
	AT 406383	T	20080915	AT 2003-763763	20030709
	ES 2304146	T3	20080916	ES 2003-763764	20030709
	EP 1995255	A1	20081126	EP 2008-14706	20030709
	R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LI, LU, MC, NL, PT, RO, SE, SI, SK, TR, LT, LV			
	ES 2311734	T3	20090216	ES 2003-763763	20030709
	US 20060167227	A1	20060727	US 2005-521140	20051031
PRAI	AT 2002-1053	A	20020712		
	EP 2003-763763	A3	20030709		
	WO 2003-EP7389	W	20030709		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
WO 2004007547	ICM	C07K014-47
	ICS	C12N015-12; G01N033-53; A01K067-027; A61K039-00; C12P021-02; C07K016-00
	IPCI	C07K0014-47 [ICM, 7]; C07K0014-435 [ICM, 7, C*]; C12N0015-12 [ICS, 7]; G01N0033-53 [ICS, 7]; A01K0067-027 [ICS, 7]; A61K0039-00 [ICS, 7]; C12P0021-02 [ICS, 7]; C07K0016-00 [ICS, 7]
	IPCR	A01K0067-027 [I, C*]; A01K0067-027 [I, A]; A61K0039-00 [N, C*]; A61K0039-00 [N, A]; C07K0014-435 [I, C*]; C07K0014-47 [I, A]; C12N0005-10 [I, C*]; C12N0005-10 [I, A]; C12N0015-09 [I, C*]; C12N0015-09 [I, A]; C12N0015-12 [I, C*]; C12N0015-12 [I, A]; C12N0015-85 [I, C*]; C12N0015-85 [I, A]; C12Q0001-02 [I, C*]; C12Q0001-02 [I, A]; C12Q0001-68 [I, C*]; C12Q0001-68 [I, A]; G01N0033-15 [I, C*]; G01N0033-15 [I, A]; G01N0033-50 [I, C*]; G01N0033-50 [I, A]; G01N0033-68 [I, C*]; G01N0033-68 [I, A]
	ECLA	C07K014/47A3; A01K067/027M4; C12N015/85A3A3; K01K; K01K; K01K; K61K; M07K; M07K
AU 2003253044	IPCI	C07K0014-47 [ICM, 7]; C07K0014-435 [ICM, 7, C*];

		A61K0039-00 [ICS,7]; C12P0021-02 [ICS,7]; C07K0016-00 [ICS,7]; C12N0015-12 [ICS,7]; G01N0033-53 [ICS,7]; A01K0067-027 [ICS,7]
	ECLA	C07K014/47A3; A01K067/027M4; C12N015/85A3A3; K01K; K01K; K01K; K61K; M07K; M07K
EP 1521774	IPCI	C07K0014-435 [I,C]; C07K0014-47 [I,A]; A01K0067-027 [I,C]; A01K0067-027 [I,A]; A61K0039-00 [I,C]; A61K0039-00 [I,A]; C07K0016-00 [I,C]; C07K0016-00 [I,A]; C12N0015-12 [I,C]; C12N0015-12 [I,A]; C12P0021-02 [I,C]; C12P0021-02 [I,A]; G01N0033-53 [I,C]; G01N0033-53 [I,A]
	IPCR	C12N0005-10 [I,C*]; C12N0005-10 [I,A]; C12N0015-09 [I,C*]; C12N0015-09 [I,A]; C12N0015-85 [I,C*]; C12N0015-85 [I,A]; C12Q0001-02 [I,C*]; C12Q0001-02 [I,A]; C12Q0001-68 [I,C*]; C12Q0001-68 [I,A]; G01N0033-15 [I,C*]; G01N0033-15 [I,A]; G01N0033-50 [I,C*]; G01N0033-50 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
	ECLA	C07K014/47A3; A01K067/027M4; C12N015/85A3A3; K01K; K01K; K01K; K61K; M07K; M07K; G01N033/50D2J4; G01N033/68V2; K01K; S01N
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	IPCR	A01K0067-027 [I,C*]; A01K0067-027 [I,A]; A61K0039-00 [N,C*]; A61K0039-00 [N,A]; C07K0014-435 [I,C*]; C07K0014-47 [I,A]; C12N0005-10 [I,C*]; C12N0005-10 [I,A]; C12N0015-09 [I,C*]; C12N0015-09 [I,A]; C12N0015-12 [I,C*]; C12N0015-12 [I,A]; C12N0015-85 [I,C*]; C12N0015-85 [I,A]; C12Q0001-02 [I,C*]; C12Q0001-02 [I,A]; C12Q0001-68 [I,C*]; C12Q0001-68 [I,A]; G01N0033-15 [I,C*]; G01N0033-15 [I,A]; G01N0033-50 [I,C*]; G01N0033-50 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
	ECLA	C07K014/47A3; A01K067/027M4; C12N015/85A3A3; K01K; K01K; K01K; K61K; M07K; M07K
JP 2006515270	IPCI	C07K0014-47 [I,A]; C07K0014-435 [I,C*]; C12N0015-09 [I,A]; C12P0021-02 [I,A]; C12Q0001-02 [I,A]; A01K0067-027 [I,A]; G01N0033-15 [I,A]; G01N0033-50 [I,A]; A61K0039-00 [I,A]; A61P0025-28 [I,A]; A61P0025-00 [I,C*]; A61K0039-39 [I,A]; C12P0021-08 [N,A]
	IPCR	C07K0014-435 [I,C]; C07K0014-47 [I,A]; A01K0067-027 [I,C]; A01K0067-027 [I,A]; A61K0039-00 [I,C]; A61K0039-00 [I,A]; A61K0039-39 [I,C]; A61K0039-39 [I,A]; A61P0025-00 [I,C]; A61P0025-28 [I,A]; C12N0005-10 [I,C*]; C12N0005-10 [I,A]; C12N0015-09 [I,C]; C12N0015-09 [I,A]; C12N0015-12 [I,C*]; C12N0015-12 [I,A]; C12N0015-85 [I,C*]; C12N0015-85 [I,A]; C12P0021-02 [I,C]; C12P0021-02 [I,A]; C12P0021-08 [N,C]; C12P0021-08 [N,A]; C12Q0001-02 [I,C]; C12Q0001-02 [I,A]; C12Q0001-68 [I,C*]; C12Q0001-68 [I,A]; G01N0033-15 [I,C]; G01N0033-15 [I,A]; G01N0033-50 [I,C]; G01N0033-50 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
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4C085/AA03; 4C085/AA38; 4C085/BB11; 4C085/CC21;
 4C085/CC28; 4C085/DD86; 4C085/FF24; 4H045/AA10;
 4H045/AA11; 4H045/AA20; 4H045/AA30; 4H045/BA10;
 4H045/CA45; 4H045/DA00; 4H045/DA76; 4H045/EA20;
 4H045/EA50; 4H045/FA72; 4H045/FA74

AT 391781 IPCI C12N0015-12 [I,C]; C12N0015-12 [I,A]; A01K0067-027
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 A61K0039-00 [N,A]; C07K0014-435 [I,C*]; C07K0014-47
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 C12N0005-10 [I,C*]; C12N0005-10 [I,A]; C12N0015-09
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AT 406383 IPCI C07K0014-435 [I,C]; C07K0014-47 [I,A]; A01K0067-027
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ES 2304146 IPCI C12N0015-12 [I,C]; C12N0015-12 [I,A]; A01K0067-027
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EP 1995255 IPCI C07K0014-47 [I,A]; C07K0014-435 [I,C*]; C12N0015-85
 [I,A]

ES 2311734 IPCI C07K0014-435 [I,C]; C07K0014-47 [I,A]; A01K0067-027
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C12P0021-02 [I,C]; C12P0021-02 [I,A]; G01N0033-53
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 G01N033/50D2J4; G01N033/68V2; K01K; K01K; K01K; K61K;
 M07K; M07K; K01K; S01N; S01N
 IPCI C07K0014-705 [I,A]; C07K0014-435 [I,C*]
 IPCR C07K0014-435 [I,C]; C07K0014-705 [I,A]
 NCL 530/350.000; 435/069.100; 435/252.300; 435/320.100;
 536/023.500
 ECLA C07K014/47A3; G01N033/50D2J4; G01N033/68V2; S01N
 AB Described are novel N- and C-terminally double truncated
 tau mols., (type IA, IB, IIA and IIB tau mols.) as well
 as methods for providing these mols., both from recombinant and biol.
 sources. Moreover, screening methods using these mols. in connection with
 Alzheimer's diagnosis and therapy are provided.
 ST tau protein fragment isolation sequence Alzheimer diagnosis
 monoclonal antibody
 IT Transformation, genetic
 (Alzheimer's disease model; N- and C-terminally truncated
 tau proteins isolation and characterization for diagnostic,
 therapeutic and Alzheimer's disease model uses)
 IT Disease models
 (Alzheimer's disease; N- and C-terminally truncated
 tau proteins isolation and characterization for diagnostic,
 therapeutic and Alzheimer's disease model uses)
 IT Vaccines
 (Alzheimer's disease; truncated Tau proteins of
 humans)
 IT Tau factor
 RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
 (Properties); BIOL (Biological study); USES (Uses)
 (N- and C-terminally double truncated; N- and C-terminally
 truncated tau proteins isolation and characterization
 for diagnostic, therapeutic and Alzheimer's disease model uses)
 IT Alzheimer's disease
 Anti-Alzheimer's agents
 Biomarkers
 Brain
 Diagnosis
 Human
 Neuron
 Oxidative stress, biological
 (N- and C-terminally truncated tau proteins
 isolation and characterization for diagnostic, therapeutic and
 Alzheimer's disease model uses)
 IT Synthetic gene
 RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
 (Uses)
 (animal; for truncated Tau proteins of humans)
 IT Microtubule
 (assembly; N- and C-terminally truncated tau
 proteins isolation and characterization for diagnostic, therapeutic and
 Alzheimer's disease model uses)
 IT Protein sequences

(for truncated Tau proteins of humans)

IT Antibodies and Immunoglobulins
 RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (monoclonal, DC44, DC82, DC136; N- and C-terminally truncated
 tau proteins isolation and characterization for diagnostic,
 therapeutic and Alzheimer's disease model uses)

IT Gene, animal
 RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
 (Uses)
 (synthetic; for truncated Tau proteins of humans)

IT Antibodies and Immunoglobulins
 RL: BSU (Biological study, unclassified); BIOL (Biological study)
 (to Alzheimer disease derived tau; N- and C-terminally
 truncated tau proteins isolation and characterization
 for diagnostic, therapeutic and Alzheimer's disease model uses)

IT Transferrins
 RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
 (Properties); BIOL (Biological study); USES (Uses)
 (τ -transferrins; N- and C-terminally truncated
 tau proteins isolation and characterization for diagnostic,
 therapeutic and Alzheimer's disease model uses)

IT 649163-53-1, 239-333-Tau factor (human type IA) 649163-54-2,
 237-333-Tau factor (human type IA) 649163-55-3, 239-318-
 Tau factor (human type IA) 649163-56-4, 239-326-Tau
 factor (human type IB) 649163-57-5, 239-328-Tau factor (human
 type IB) 649163-58-6, 239-331-Tau factor (human type IB)
 649163-59-7, 239-334-Tau factor (human type IB) 649163-60-0,
 239-340-Tau factor (human type IB) 649163-61-1, 239-343-
 Tau factor (human type IB) 649163-62-2, 208-302-Tau
 factor (human type IB) 649163-63-3, 69-333-Tau factor (human
 type IIA) 649163-64-4, 93-333-Tau factor (human type IIA)
 649163-65-5, 69-363-Tau factor (human type IIA) 649163-66-6,
 93-363-Tau factor (human type IIA) 649163-67-7, 93-302-
 Tau factor (human type IIA) 649163-68-8, 69-302-Tau
 factor (human type IIA) 649163-69-9, 93-332-Tau factor (human
 type IIA) 649163-70-2, 69-332-Tau factor (human type IIA)
 649163-71-3, 6-378-Tau factor (human type IIB) 649163-72-4,
 6-347-Tau factor (human type IIB) 649163-73-5, 268-333-
 Tau factor (human) 649163-74-6, 248-333-Tau factor
 (human) 649163-75-7, 258-333-Tau factor (human) 649163-76-8,
 263-333-Tau factor (human) 649163-77-9, (239-247)-(263-333)-
 Tau factor (human) 649163-78-0, (239-255)-(263-333)-Tau
 factor (human) 649163-79-1, (239-262)-(268-333)-Tau factor
 (human)
 RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
 (Biological study)
 (amino acid sequence; N- and C-terminally truncated
 tau proteins isolation and characterization for diagnostic,
 therapeutic and Alzheimer's disease model uses)

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
 RE

- (1) Anon; WO 0118546 A2 CAPLUS
- (2) Anon; WO 0165252 A1 CAPLUS
- (3) Anon; WO 02055720 A2 CAPLUS
- (4) Anon; WO 02059150 A2 CAPLUS
- (5) Anon; WO 02062851 A1 CAPLUS
- (6) Anon; WO 9630766 A1 CAPLUS

=> file registry
 COST IN U.S. DOLLARS

SINCE FILE TOTAL

FULL ESTIMATED COST	ENTRY 24.70	SESSION 25.36
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
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649163-68-8/rn or 649163-69-9/rn or 649163-70-2/rn or 649163-71-3/rn
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      1 649163-68-8/RN
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      1 649163-70-2/RN
      1 649163-71-3/RN
L4    8 649163-64-4/RN OR 649163-65-5/RN OR 649163-66-6/RN OR 649163-67
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      649163-71-3/RN
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	ENTRY	SESSION
FULL ESTIMATED COST	4.32	29.68
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
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FILE LAST UPDATED: 27 Mar 2009 (20090327/ED)

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=> s l4

L5 1 L4

=> file uspatfull

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.50	30.18
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
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FILE COVERS 1971 TO PATENT PUBLICATION DATE: 26 Mar 2009 (20090326/PD)
FILE LAST UPDATED: 26 Mar 2009 (20090326/ED)
HIGHEST GRANTED PATENT NUMBER: US7509687
HIGHEST APPLICATION PUBLICATION NUMBER: US20090083889
CA INDEXING IS CURRENT THROUGH 26 Mar 2009 (20090326/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 26 Mar 2009 (20090326/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2008
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2008

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L6 0 L4

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L1 48 S KONTSEKOVA E?/AU
L2 10 S L1 AND TAU
L3 4 S L2 AND TRUNCATED

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	1.69	31.87
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
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L5 FILE 'CAPLUS' ENTERED AT 15:11:11 ON 28 MAR 2009
1 S L4

L6 FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009
0 S L4

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=> d ibib abs 15 1

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2004:60551 CAPLUS
DOCUMENT NUMBER: 140:124832
TITLE: Truncated tau proteins
INVENTOR(S): Kontsekova, Eva
PATENT ASSIGNEE(S): Axon Neuroscience Forschungs- und Entwicklungs GmbH,
Austria
SOURCE: PCT Int. Appl., 97 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004007547	A2	20040122	WO 2003-EP7389	20030709
WO 2004007547	A3	20040722		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2003253044	A1	20040202	AU 2003-253044	20030709
EP 1521774	A2	20050413	EP 2003-763763	20030709
EP 1521774	B1	20080827		
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
CN 1668641	A	20050914	CN 2003-816647	20030709
JP 2006515270	T	20060525	JP 2004-520541	20030709
AT 391781	T	20080415	AT 2003-763764	20030709
AT 406383	T	20080915	AT 2003-763763	20030709
ES 2304146	T3	20080916	ES 2003-763764	20030709
EP 1995255	A1	20081126	EP 2008-14706	20030709
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LI, LU, MC, NL, PT, RO, SE, SI, SK, TR, LT, LV			
ES 2311734	T3	20090216	ES 2003-763763	20030709
US 20060167227	A1	20060727	US 2005-521140	20051031
PRIORITY APPLN. INFO.:			AT 2002-1053	A 20020712
			EP 2003-763763	A3 20030709
			WO 2003-EP7389	W 20030709
AB	Described are novel N- and C-terminally double truncated tau mols., (type IA, IB, IIA and IIB tau mols.) as well as methods for providing these mols., both from recombinant and biol. sources. Moreover, screening methods using these mols. in connection with Alzheimer's diagnosis and			

therapy are provided.
REFERENCE COUNT: 6

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L5 1 S L4

FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009

L6 0 S L4

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MOST RECENT UPDATE: 200918 <200918/DW>
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ECLA, F-Term and FI-Term classifications are complete
to the end of 2008.
No update date (UP) has been created for the reclassified
documents, but they can be identified by
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L3 4 S L2 AND TRUNCATED
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FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009

L6 0 S L4

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0 649163-68-8/RN
0 649163-69-9/RN
0 649163-70-2/RN
0 649163-71-3/RN
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649163-71-3/RN

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649163-71-3/RN

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COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

5.87

42.24

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

	ENTRY	SESSION
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L5 1 S L4

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L6 0 S L4

FILE 'CAPLUS' ENTERED AT 15:11:57 ON 28 MAR 2009

FILE 'WPIDS' ENTERED AT 15:13:48 ON 28 MAR 2009

L7 0 S L4
L8 0 S L5

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=> d l4 1

L4 ANSWER 1 OF 8 REGISTRY COPYRIGHT 2009 ACS on STN
RN 649163-71-3 REGISTRY
ED Entered STN: 11 Feb 2004
CN 6-378-Tau factor (human type IIB) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 19: PN: WO2004007547 SEQID: 19 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L4 ANSWER 4 OF 8 REGISTRY COPYRIGHT 2009 ACS on STN
RN 649163-68-8 REGISTRY
ED Entered STN: 11 Feb 2004
CN 69-302-Tau factor (human type IIA) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 16: PN: WO2004007547 SEQID: 16 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L4 ANSWER 8 OF 8 REGISTRY COPYRIGHT 2009 ACS on STN
RN 649163-64-4 REGISTRY
ED Entered STN: 11 Feb 2004
CN 93-333-Tau factor (human type IIA) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 12: PN: WO2004007547 SEQID: 12 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

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*** USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE ***
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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L2 10 S L1 AND TAU

L3 4 S L2 AND TRUNCATED
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L5 1 S L4

FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009
L6 0 S L4

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L7 0 S L4
L8 0 S L5

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 649163-69-9/RN OR 649163-70-2/RN OR 649163-71-3/RN

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